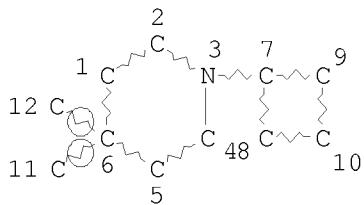


=> d 11  
L1 HAS NO ANSWERS  
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NODE ATTRIBUTES:  
DEFAULT MLEVEL IS ATOM  
DEFAULT ECLEVEL IS LIMITED

## GRAPH ATTRIBUTES:

RSPEC 8 3  
NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

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FULL SEARCH INITIATED 12:08:53 FILE 'REGISTRY'
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L3 93 SEA SSS FUL L1

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COST IN U.S. DOLLARS  
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FILE 'CAPLUS' ENTERED AT 12:08:57 ON 17 SEP 2008  
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FILE COVERS 1907 - 17 Sep 2008 VOL 149 ISS 12  
FILE LAST UPDATED: 16 Sep 2008 (20080916/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply.

They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>

=> s 13  
L4 12 L3

=> d bib 1-12

L4 ANSWER 1 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2008:353313 CAPLUS  
DN 148:379624  
TI Benzimidazolecarboxamide derivatives as bradykinin B1 receptor modulators and their preparation, pharmaceutical compositions and use in the treatment of inflammation and pain  
IN Chandrasekhar, Jayaraman; Guo, Qin; Ihle, David C.; Ge, Ping; Wustrow, David J.; Chenard, Bertrand L.; Hodgetts, Kevin J.  
PA Neurogen Corporation, USA  
SO PCT Int. Appl., 80pp.  
CODEN: PIXXD2  
DT Patent  
LA English  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2008033739	A2	20080320	WO 2007-US77959	20070910
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
PRAI	US 2006-825308P	P	20060912		
OS	MARPAT	148:379624			

L4 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2007:1392027 CAPLUS  
DN 148:54908  
TI Preparation of spirocyclic sulfonamides and related compounds as modulators of bradykinin receptor activity  
IN Hodgetts, Kevin J.; Ihle, David C.; Li, Guiying; Ge, Ping; Chenard, Bertrand L.; Wustrow, David J.  
PA Neurogen Corporation, USA  
SO PCT Int. Appl., 82pp.  
CODEN: PIXXD2

DT Patent  
LA English  
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2007140383	A2	20071206	WO 2007-US69918	20070530
	WO 2007140383	A3	20080124	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG,	

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 PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN,  
 TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW  
 RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,  
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 BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW,  
 GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,  
 BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRAI US 2006-803419P P 20060530  
 OS MARPAT 148:54908

L4 ANSWER 3 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:1215844 CAPLUS  
 DN 147:486464  
 TI Preparation of 3-(piperazin-1-yl)cyclobut-3-ene-1,2-dione derivatives as  
 antiviral agents  
 IN Bachand, Carol; Deon, Daniel H.; Ruediger, Edward H.  
 PA Bristol-Myers Squibb Company, USA  
 SO U.S. Pat. Appl. Publ., 38pp.  
 CODEN: USXXCO

DT Patent  
 LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20070249624	A1	20071025	US 2007-737354	20070419
	WO 2007127731	A1	20071108	WO 2007-US67302	20070424
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				

PRAI US 2006-794699P P 20060425  
 OS MARPAT 147:486464

L4 ANSWER 4 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2007:563421 CAPLUS  
 DN 147:9912  
 TI Preparation of diazacycloalkane-containing N,N-bis(1H-imidazol-2-ylmethyl)amines and related N,N-bis(heteroaryl methyl)amines as antagonists of chemokine receptor CXCR4  
 IN Kokubo, Masaya; Ochiai, Hiroshi; Takaoka, Yoshikazu; Shibayama, Shiro  
 PA Ono Pharmaceutical Co., Ltd., Japan  
 SO PCT Int. Appl., 332pp.  
 CODEN: PIXXD2

DT Patent  
 LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2007058322	A1	20070524	WO 2006-JP323015	20061117
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,				

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MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO,  
RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT,  
TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW  
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,  
IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,  
CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,  
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,  
KG, KZ, MD, RU, TJ, TM

EP 1961744 A1 20080827 EP 2006-832893 20061117  
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,  
IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR

PRAI JP 2005-334937 A 20051118  
JP 2006-49378 A 20060224  
WO 2006-JP323015 W 20061117

OS MARPAT 147:9912

RE.CNT 42 THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2007:203199 CAPLUS  
DN 146:274233  
TI Preparation of spiro-fused piperidines as modulators of muscarinic receptors  
IN Makings, Lewis R.; Garcia-Guzman Blanco, Miguel; Hurley, Dennis J.; Drutu, Ioana; Raffai, Gabriel; Bergeron, Daniele M.; Nakatani, Akiko; Termin, Andreas P.; Silina, Alina  
PA USA  
SO U.S. Pat. Appl. Publ., 500pp., Cont.-in-part of U.S. Ser. No. 208,386.  
CODEN: USXXCO  
DT Patent  
LA English  
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20070043023	A1	20070222	US 2006-359960	20060222
	US 20060079505	A1	20060413	US 2005-208386	20050819
PRAI	US 2004-602731P	P	20040819		
	US 2005-208386	A2	20050819		

OS MARPAT 146:274233

L4 ANSWER 6 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
AN 2006:693858 CAPLUS  
DN 145:201982  
TI 4-Substituted-8-(1-phenyl-cyclohexyl)-2,8-diaza-spiro[4.5]decan-1-one as a novel class of highly selective GlyT1 inhibitors with superior pharmacological and pharmacokinetic parameters  
AU Alberati, Daniela; Hainzl, Dominik; Jolidon, Synese; Kurt, Anke; Pinard, Emmanuel; Thomas, Andrew W.; Zimmerli, Daniel  
CS Discovery Biology, Pharmaceuticals Division, F. Hoffmann-La Roche Ltd., Basel, CH-4070, Switz.  
SO Bioorganic & Medicinal Chemistry Letters (2006), 16(16), 4321-4325  
CODEN: BMCLE8; ISSN: 0960-894X  
PB Elsevier B.V.  
DT Journal  
LA English  
OS CASREACT 145:201982  
RE.CNT 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD  
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 7 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2006:193777 CAPLUS  
 DN 144:274274  
 TI Preparation of basic group-containing heterocyclic compounds as CXCR4 antagonists  
 IN Kokubo, Masaya; Takaoka, Yoshikazu; Shibayama, Shiro  
 PA Ono Pharmaceutical Co., Ltd, Japan  
 SO PCT Int. Appl., 305 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA Japanese  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006022454	A1	20060302	WO 2005-JP16066	20050826
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	EP 1790639	A1	20070530	EP 2005-776646	20050826
	R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
	US 20080009495	A1	20080110	US 2007-661270	20070227
PRAI	JP 2004-248431	A	20040827		
	JP 2005-100039	A	20050330		
	JP 2005-190741	A	20050629		
	WO 2005-JP16066	W	20050826		
OS	MARPAT 144:274274				
RE.CNT 15	THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD				
	ALL CITATIONS AVAILABLE IN THE RE FORMAT				

L4 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2006:15778 CAPLUS  
 DN 144:108321  
 TI Preparation of amino cyclopentyl heterocyclic and carbocyclic modulators of chemokine receptor activity  
 IN Yang, Lihu; Lin, Songnian; Morriello, Gregori; Guo, Liangqin; Zhou, Changyou  
 PA Merck & Co., Inc., USA  
 SO PCT Int. Appl., 141 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2006001958	A2	20060105	WO 2005-US17836	20050520
	WO 2006001958	A3	20060824		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,				

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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
AU 2005257859	A1 20060105	AU 2005-257859	20050520
CA 2567851	A1 20060105	CA 2005-2567851	20050520
EP 1753740	A2 20070221	EP 2005-785401	20050520
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, LV, MK, YU			
CN 1956975	A 20070502	CN 2005-80016231	20050520
JP 2008502719	T 20080131	JP 2007-527495	20050520
IN 2006DN06362	A 20070831	IN 2006-DN6362	20061030
PRAI US 2004-573625P	P 20040521		
WO 2005-US17836	W 20050520		
OS MARPAT 144:108321			

L4 ANSWER 9 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2005:612285 CAPLUS  
 DN 143:133293  
 TI Preparation of spiroindoline and spiroisoquinoline compounds as Mas receptor ligands  
 IN Boatman, Douglas P.; Adams, John W.; Moody, Jeanne V.; Babych, Eric D.; Schrader, Thomas O.  
 PA Arena Pharmaceuticals, Inc., USA  
 SO PCT Int. Appl., 224 pp.  
 CODEN: PIXXD2

DT Patent  
 LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2005063745	A2	20050714	WO 2004-US43609	20041222
	WO 2005063745	A3	20060316		
	WO 2005063745	A9	20070201		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, SM RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2004309419	A1	20050714	AU 2004-309419	20041222
	CA 2546147	A1	20050714	CA 2004-2546147	20041222
	EP 1716148	A2	20061102	EP 2004-815636	20041222
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, BA, HR, IS, YU				
	CN 1972944	A	20070530	CN 2004-80038166	20041222
	JP 2007516298	T	20070621	JP 2006-547461	20041222
	IN 2006KN02015	A	20070518	IN 2006-KN2015	20060718
	US 20070254903	A1	20071101	US 2007-583839	20070308
PRAI	US 2003-532546P	P	20031223		
	US 2004-539554P	P	20040126		

US 2004-565251P P 20040423  
 WO 2004-US43609 W 20041222  
 OS MARPAT 143:133293

L4 ANSWER 10 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2004:1124588 CAPLUS  
 DN 142:69197  
 TI CCR-2 antagonists for treatment of neuropathic pain  
 IN Abbadie, Catherine; Lindia, Jill Ann; Wang, Hao  
 PA Merck & Co., Inc., USA  
 SO PCT Int. Appl., 304 pp.  
 CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004110376	A2	20041223	WO 2004-US17499	20040602
	WO 2004110376	A3	20050224		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 20060205761	A1	20060914	US 2005-559701	20051206
PRAI	US 2003-476391P	P	20030606		
	US 2003-531637P	P	20031222		
	WO 2004-US17499	W	20040602		
OS	MARPAT 142:69197				

L4 ANSWER 11 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2004:802715 CAPLUS

DN 141:314157

TI Preparation of amino cyclobutylamide modulators of chemokine receptor activity

IN Jiao, Richard; Yang, Lihu

PA Merck & Co. Inc., USA

SO PCT Int. Appl., 108 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004082682	A1	20040930	WO 2004-US7792	20040315
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,				

TD, TG  
 AU 2004222336 A1 20040930 AU 2004-222336 20040315  
 CA 2519220 A1 20040930 CA 2004-2519220 20040315  
 EP 1617841 A1 20060125 EP 2004-720791 20040315  
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK  
 CN 1787818 A 20060614 CN 2004-80013143 20040315  
 JP 2006520783 T 20060914 JP 2006-507176 20040315  
 IN 2005DN03929 A 20070824 IN 2005-DN3929 20050902  
 US 20060211722 A1 20060921 US 2005-549739 20050919  
 PRAI US 2003-456047P P 20030318  
 WO 2004-US7792 A 20040315  
 OS MARPAT 141:314157

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD  
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 AN 2001:247141 CAPLUS  
 DN 134:266327  
 TI Preparation of spiro[bicyclic-azacycloalkyl and -cycloalkyl] derivatives  
 and uses as alpha 1a adrenergic receptor antagonists  
 IN Hoffman, Jacob M.  
 PA Merck & Co., Inc., USA  
 SO PCT Int. Appl., 140 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

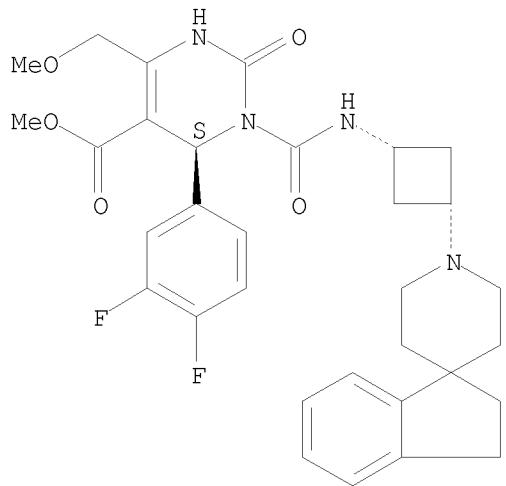
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001022919	A2	20010405	WO 2000-US26387	20000926
	WO 2001022919	A3	20020711		
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	AU 2000077166	A	20010430	AU 2000-77166	20000926
PRAI	US 1999-157207P	P	19990930		
	WO 2000-US26387	W	20000926		
OS	MARPAT 134:266327				

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L4 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2008 ACS on STN  
 IT 332187-08-3P 332187-09-4P 332187-37-8P  
 332187-38-9P  
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological  
 study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);  
 BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (preparation of spirobicyclicazacycloalkyl and cycloalkyl derivs. and uses  
 as alpha a adrenergic receptor antagonists)  
 RN 332187-08-3 CAPLUS  
 CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[cis-3-(2,3-  
 dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-  
 1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, hydrochloride

(1:1), (6S)- (CA INDEX NAME)

Absolute stereochemistry.

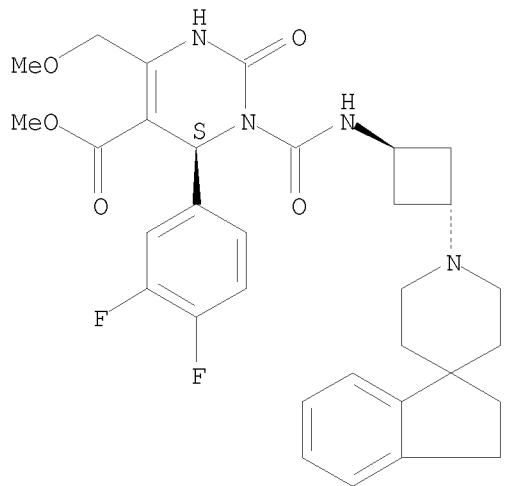


● HCl1

RN 332187-09-4 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[trans-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, hydrochloride (1:1), (6S)- (CA INDEX NAME)

Absolute stereochemistry.

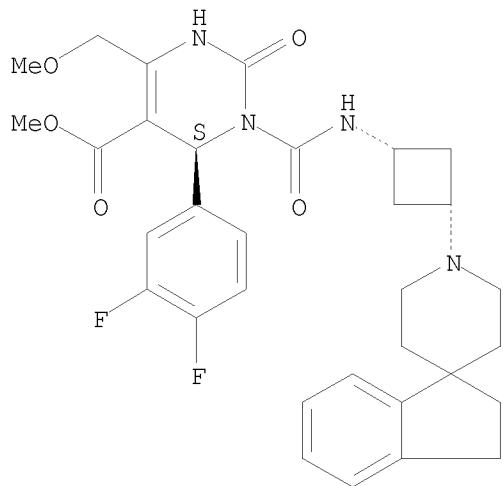


● HCl1

RN 332187-37-8 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[cis-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, (6S)- (CA INDEX NAME)

Absolute stereochemistry.



RN 332187-38-9 CAPLUS

CN 5-Pyrimidinecarboxylic acid, 6-(3,4-difluorophenyl)-1-[[[trans-3-(2,3-dihydrospiro[1H-indene-1,4'-piperidin]-1'-yl)cyclobutyl]amino]carbonyl]-1,2,3,6-tetrahydro-4-(methoxymethyl)-2-oxo-, methyl ester, (6S)- (CA INDEX NAME)

Absolute stereochemistry.

